

Prisca 5.0.2.37
Date of report: 14-09-2017

Patient data			
Name	MRS. SIMPLE	Patient ID	1709220325/AMB
Birthdate	15-02-1988	Sample ID	1709220325/AMB
Age at sample date	29.6	Sample Date	13-09-2017
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	46	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies			no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	8.67 mIU/ml	1.63	Gestational age 12 + 4
fb-hCG	55.5 ng/ml	1.52	Method CRL Robinson
			Scan date 11-09-2017
Risks at sampling date			Crown rump length in mm 63.5
Age risk		1:687	Nuchal translucency MoM 1.23
Biochemical T21 risk		1:4169	Nasal bone present
Combined trisomy 21 risk		1:7980	Sonographer .
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT MD
			<p>Trisomy 21</p> <p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 7980 women with the same data, there is one woman with a trisomy 21 pregnancy and 7979 women with not affected pregnancies.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

